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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/849,374	05/19/2004	Bjorn Olstad	15-DS-00549 (MHM 13109US0	6522
23446 OSSIJA008 OSSIJA008 OSSIJA008 OSSIJA008 OSSIJA008 OSSIJA00 O			EXAMINER	
			CHAO, ELMER M	
CHICAGO, IL	.60661		ART UNIT	PAPER NUMBER
			3737	
			MAIL DATE	DELIVERY MODE
			05/01/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Application No. Applicant(s) 10/849,374 OLSTAD, BJORN Office Action Summary Art Unit Examiner ELMER CHAO 3737 -- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --Period for Reply A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS. WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b). Status 1) Responsive to communication(s) filed on 19 December 2007. 2a) This action is FINAL. 2b) This action is non-final. 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213. Disposition of Claims 4) Claim(s) 1-21 is/are pending in the application. 4a) Of the above claim(s) _____ is/are withdrawn from consideration. 5) Claim(s) _____ is/are allowed. 6) Claim(s) 1-21 is/are rejected. 7) Claim(s) _____ is/are objected to. 8) Claim(s) _____ are subject to restriction and/or election requirement. Application Papers 9) The specification is objected to by the Examiner. 10) The drawing(s) filed on is/are; a) accepted or b) objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abevance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d). 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152. Priority under 35 U.S.C. § 119 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received.

1) Notice of References Cited (PTO-892)

Notice of Draftsperson's Patent Drawing Review (PTO-948)

Information Disclosure Statement(s) (FTO/S5/08)
 Paper No(s)/Mail Date _______.

Attachment(s)

Interview Summary (PTO-413)
 Paper No(s)/Mail Date.

6) Other:

5 Notice of Informal Patent Application

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DETAILED ACTION

1. Acknowledgement is made of the amendment filed 12/19/2007.

Response to Arguments

 Applicant's arguments filed 12/19/2007 have been fully considered but they are not persuasive.

Regarding Applicant's arguments with respect to newly amended claims 1 and 13, Examiner asserts that all of the limitations are still read on by Roundhill et al. Roundhill et al. teach presetting values in an analytical instrument using said clinically relevant information (col. 10, lines 13-15). Specifically, the clinically relevant information would be in this case the "control points beneath the border". These points are then Doppler processed so inherently an analytical instrument would have to be preset with those values so that the information can be processed by the analytical instrument running the Doppler process.

Regarding claims 8-10, a new ground of rejection has been provided as necessitated by the amendments.

Double Patenting

3. The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., In re Berg, 140 F.3d 1428, 46 USPQ2d 1226 (Fed.

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Cir. 1998); In re Goodman, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); In re Longi, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); In re Van Omum, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); In re Vogel, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and In re Thorington, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1,321(c) or 1,321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

4. Claims 1, 3, 4, 5, and 7 are provisionally rejected on the ground of nonstatutory double patenting over claims 6, 9, and 11-16 of copending Application No. 11/082,540. This is a provisional double patenting rejection since the conflicting claims have not yet been patented.

The subject matter claimed in the instant application is fully disclosed in the referenced copending application and would be covered by any patent granted on that copending application since the referenced copending application and the instant application are claiming common subject matter, as follows: a method of using an ultrasound machine including locating at least one anatomical landmark; locating at least one clinically relevant location based on the at least one anatomical landmark; extracting clinically relevant information from said at least one clinically relevant location within the heart; and displaying indicia overlaying said at least one clinically relevant location.

Furthermore, there is no apparent reason why applicant would be prevented from presenting claims corresponding to those of the instant Application/Control Number: 10/849,374

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application in the other copending application. See *In re Schneller*, 397 F.2d 350, 158 USPQ 210 (CCPA 1968). See also MPEP § 804.

5. Claims 1-5, 7, 10, and 11 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-11 of copending Application No. 10/848,721. Although the conflicting claims are not identical, they are not patentably distinct from each other because the conflicting claims are narrower in scope than the present application claims.

This is a <u>provisional</u> obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35
 U.S.C. 102 that form the basis for the rejections under this section made in this
 Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- Claims 1-21 are rejected under 35 U.S.C. 102(b) as being anticipated by Roundhill et al. (U.S. 6,447,453 B1; also see Publication U.S. 2002/0072672 A1).

Regarding **claims 1-4**, **and 7**, Roundhill et al. teach a method for using an ultrasound machine comprising locating at least one anatomical landmark within the cardiac structure and generating position information of said at least one

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anatomical landmark comprising the apex of the heart (Fig. 4); locating at least one clinically relevant location within the heart based on said position information of said at least one anatomical landmark (Fig. 8, see the border of the LV); and extracting clinically relevant information from said at least one clinically relevant location within the heart, wherein the clinically relevant information can comprise Doppler profile information (col. 10, lines 12-15), wherein the extracting involves presetting at least one m-mode, Doppler sample volume, and region-of-interest (col. 9, line 62 – col. 10, line 51); and displaying indicia overlaying said at least one clinically relevant location on a display of said ultrasound machine (Fig. 5, Items 26, 36, & 46; Fig. 11, Item 100).

Regarding **claims 8-10**, Roundhill et al. teach the preset values being m-mode, Doppler sample volume, or a ROI (col. 9, line 62 – col. 10, line 10, "m-mode"; col. 10, lines 10-20).

Regarding claim 5, Roundhill et al. teach the clinically relevant location being the LV portion of the heart (see Fig. 13a).

Regarding claim 6, Roundhill et al. teach that locating at least one clinically relevant location includes performing edge detection of said at least one myocardial segment of the heart to locate endocardium of said at least one myocardial segment (see Fig. 12; claim 12; col. 11, line 58 – col. 12, line 10).

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Regarding **claims 11 and 12**, Roundhill et al. teach tracking the apex and control points as the heart is beating and automatically drawing cardiac borders (col. 10, lines 52 – 67; col. 13, lines 12-31).

Regarding **claims 13-21**, Roundhill et al. teach the method steps as discussed above which would necessitate the presence of the corresponding apparatus parts used in the method steps.

Claim Rejections - 35 USC § 103

- The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior at are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 9. Claims 8-10 are alternatively rejected under 35 U.S.C. 103(a) as being unpatentable over Roundhill et al. Roundhill et al. teach the limitations as discussed above but may not explicitly teach presetting values of M-mode and volume. However, it is well known in the art to use anatomical information to preset ultrasound imaging systems and processors. Therefore, it would have been obvious to a person of ordinary skill in the art at the time of the invention to preset values of M-mode and volume based on the anatomical scans.

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Conclusion

10. Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, THIS ACTION IS MADE FINAL.
See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

11. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Elmer Chao whose telephone number is (571)272-0674. The examiner can normally be reached on 9am-4pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Brian Casler can be reached on (571)272-4956. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Brian L Casler/ Supervisory Patent Examiner, Art Unit 3737

/E. C./ Examiner, Art Unit 3737 4/25/2008